

Ansell Perry  
1875 Harsh Avenue SE  
Massillon, Ohio 44646  
Telephone: 330-833-2811  
Fax: 330-833-6501

**JUL 27 2000***K002192*

[1] Summary

[2] Ansell Perry Inc.  
1875 Harsh Avenue SE  
Massillon, Ohio 44646

Contact: James R. Chatterton  
Telephone: 330-833-2811  
Fax: 330-833-6501

July 17, 2000

- [3] Trade Name: Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less)  
Common Name: Surgical Gloves  
Classification Name: Surgeon's Glove
- [4] Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less) meet all of the requirements of ASTM D 3577-00, Type 1.
- [5] Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less) meet all the current specifications for ASTM D 3577-00 Rubber Surgical Gloves.
- [6] Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less) are sterile disposable devices intended to be worn by operating room personnel to protect a surgical wound from contamination.
- [7] Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less) are summarized with the following technological characteristics compared to ASTM or equivalent standards.

Characteristics	Standard
Dimensions	Meets ASTM D 3577-00
Physical Properties	Meets ASTM D 3577-00, Type 1
Freedom from holes	Meets ASTM D 3577-00 • Meets ASTM D 5151-92
Powder-Free	Meets ASTM D 3577-00 Not more than 2 mg residue by mass per glove.
Protein Label Claim	This latex glove contains 50 micrograms of total water extractable protein per gram.

R002192

Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content  
Labeling Claim (50 micrograms or less) (Modification)

Ansell Perry  
1875 Harsh Avenue SE  
Massillon, Ohio 44646  
Telephone: 330-833-2811  
Fax: 330-833-6501

Biocompatibility

Primary Skin Irritation in Rabbits  
Guinea Pig Sensitization

Passes  
Passes

- [8] The performance test data of the non clinical tests are the same as mentioned immediately above.
- [9] Clinical data is not needed for medical gloves or for most devices cleared by the 510(k) process.
- [10] It is concluded that the Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less) (Modified) are as safe, as effective, and perform as well as the glove performance standards referenced above and therefore meet:

ASTM listed standards,  
FDA hole requirements, and  
labeling claims for the product.

- [11] This summary will include any other information reasonably deemed necessary by the FDA.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

JUL 27 2000

Mr. James R. Chatterton  
Vice President Regulatory  
Ansell Healthcare Products, Incorporated  
1875 Harsh Avenue South East  
Massillon, Ohio 44646

Re: K002192  
Trade Name: Dermaguard PF Brown Powder Free Latex  
Surgical Gloves; MOD  
Regulatory Class: I  
Product Code: KGO  
Dated: July 17, 2000  
Received: July 20, 2000

Dear Mr. Chatterton:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

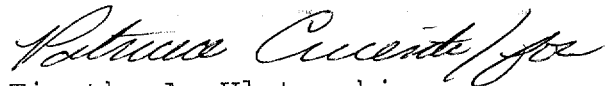
If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

Page 2 - Mr. Chatterton

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsma/dsmamain.html>".

Sincerely yours,



Timothy A. Ulatowski  
Director

Division of Dental, Infection Control  
and General Hospital Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

Attachment 2

Indications for Use Statement

510(k)  
Number  
(if known)

K002192

Device Name

Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less)

Indications for Use

Dermaguard PF Brown Powder Free Latex Surgical Gloves with Protein Content Labeling Claim (50 micrograms or less) intended use is to be worn by operating room personnel to protect a surgical wound from contamination.

PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF  
NEEDED

Concurrence of CDRH Office of Device Evaluation (ODE)



(Division Sign-Off)

Division of Dental, Infection Control  
and General Hospital

510(k) Number

OR

K002192

Prescription Use \_\_\_\_\_  
Per 21 CFR 801.109

Over-The-Counter Use

X